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CLAIMS

1. A method of treating neurodegenerative disease in a mammal comprising the steps of introducing a therapeutic effective amount of a chaperone or chaperone-like-compound into the neurological system of the mammal.
2. The method of claim 1, wherein the introducing step includes introducing the chaperone or chaperone-like-compound into the mammal by gene therapy.
3. The method of claim 1, wherein the introducing step includes directly injecting the chaperone or chaperone-like-compound into the mammal.
4. A method for screening for a test compound for chaperone-like activity for the treatment of neurodegenerative diseases comprising the steps of:
  - introducing the test compound into transfected cells in tissue culture, wherein such transfected cells produce protein aggregate; and
  - measuring the quantity of protein aggregate, wherein a test compound which decreases the quantity of protein aggregate as compared to control cells has chaperone activity.
5. A method for screening for a test compound for chaperone-like activity for the treatment of neurodegenerative diseases comprising the steps of:
  - introducing the test compound into an animal which models neurodegenerative disease;

allowing said animal to develop; and subsequently measuring the quantity of aggregates in said animal wherein decreased aggregate formation over control animals indicates chaperone-like activity.

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6. A method of treating neurodegenerative disease in a mammal comprising the step of introducing a therapeutically effective amount of a compound into said mammal wherein said compound increases the effective concentration of a chaperone in the neurological system.

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7. A method of treating neurodegenerative disease in a mammal comprising the step of introducing a therapeutically effective amount of a compound into said mammal wherein said compound increases the effective concentration or enhances the activity of a proteasome in the neurological system.

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8. A method for screening for a test compound which increases proteasome activity for the treatment of neurodegenerative diseases comprising the steps of:  
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introducing the test compound into transfected cells in tissue culture, wherein such transfected cells produce protein aggregate; and  
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measuring the quantity of protein aggregate, wherein a test compound which decreases the quantity of protein aggregate is selected.

9. A method for screening for a test compound which increases proteasome activity for the treatment of neurodegenerative diseases comprising the steps of:  
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introducing the test compound into an animal  
which models neurodegenerative disease;

allowing said animal to develop; and

subsequently measuring the quantity of

5 aggregates in said animal wherein a compound which  
shows decreased aggregate formation over control  
animals is selected.

10. Transgenic mice capable of overexpression of  
HDJ-2.